

Amendments to the Claims

1. (Currently amended) A method of treating Use of papaverine-like vasodilator for the production of a pharmaceutical composition for the treatment of an ophthalmological dysfunction dysfunctions which is are linked to circulatory disturbances of the eye or which can are to be attributed to circulatory disturbances of the eye, comprising administering a pharmaceutical composition comprising a papaverine-like vasodialator, wherein said the pharmaceutical composition is administered to be applied topically to said the eye.
2. (Currently amended) The method according to Use as set forth in claim 1 wherein characterised in that the papaverine-like vasodilator is selected from the group consisting of whichrconsists of papaverine, ethaverine, moxaverine, elziverine, their pharmacologically compatible salts and mixtures thereof.
3. (Currently amended) The method according to claim 1 Use as set forth in one of the preceding claims characterised in that wherein the ophthalmological dysfunction dysfunctions is are selected from the group consisting of which consists of glaucoma and an ophthalmological dysfunction dysfunctions linked to diabetes, for example neovascularisation glaucoma, and diabetic retinopathy.

4. (Currently amended) The method according to claim 1 Use as set forth in one of claims 1 through 3 characterised in that wherein the pharmaceutical composition is in a the form selected from the group consisting of eye drops, eye ointments, eye spray, eye tablet, gel, suspension, emulsion, powder and or granules.
5. (Currently amended) The method according to claim 1 Use as set forth in one of the preceding claims characterised in that wherein the pharmaceutical composition additionally includes a viscosity regulator, wherein the viscosity regulator has a viscosity-increasing action.
6. (Currently amended) The method according to Use as set forth in claim 5 characterised in that wherein the viscosity regulator is selected from the group which consists consisting of chondroitin sulfate, polyacrylamide, polyacrylic acid, polyacrylic resins, polyethylene glycol, cellulose derivatives, polyvinyl alcohol, polyvinyl pyrrolidone, hyaluronic acid, hyaluronates and mixtures thereof.
7. (Currently amended) A pharmaceutical composition which includes a papaverine-like vasodilator and a pharmacologically compatible viscosity regulator, wherein the papaverine-like vasodilator is selected from the group which consists consisting of moxaverine, its pharmacologically compatible salts and mixtures

thereof, and wherein the viscosity regulator is selected from the group which consists consisting of chondroitin sulfate, polyacrylamide, polyacrylic acid, polyacrylic resins, polyethylene glycol, cellulose derivatives, polyvinyl alcohol, polyvinyl pyrrolidone, hyaluronic acid, hyaluronates and mixtures thereof.

8. (Currently amended) A pharmaceutical composition as set forth in claim 7 wherein characterised in that the pharmaceutical composition is in a the form selected from the group consisting of eye drops, eye ointments, eye spray, eye tablet, gel, suspension, emulsion, powder and or granules.
9. (New) The method according to claim 3 wherein said ophthalmological dysfunction linked to diabetes is selected from the group consisting of neovascularisation glaucoma and diabetic retinopathy.